



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : A61L 2/20, 11/00 // 101/10, 101/36	A1	(11) International Publication Number: WO 00/02595 (43) International Publication Date: 20 January 2000 (20.01.00)
(21) International Application Number: PCT/EP99/04723 (22) International Filing Date: 5 July 1999 (05.07.99) (30) Priority Data: 09/113,699 10 July 1998 (10.07.98) US (71) Applicant (for all designated States except US): BOX 03 INTERNATIONAL [CH/CH]; Hintere Dorfasse 9, CH-3073 Gümligen (CH). (72) Inventor; and (75) Inventor/Applicant (for US only): DUROSELLE, Patrick [FR/FR]; 18 Rue Nungesser et Coli, F-69008 Lyon (FR). (74) Agent: BOVARD LTD.; Optingenstrasse 16, CH-3000 Bern 25 (CH).		(81) Designated States: AE, AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: A METHOD FOR DISINFECTING AND STERILIZING MICROBIAL CONTAMINATED MATERIALS (57) Abstract <p>The method for disinfecting and sterilizing microbial contaminated or infectious materials, e.g. medical instruments, waste from medical laboratories or hospitals, comprises the following steps: (a) loading the material for disinfecting and sterilizing into a vacuum-proof sterilization chamber; (b) introducing liquid hydrogen peroxide into the sterilization chamber so as to penetrate the material; (c) introducing liquid acetic acid into the sterilization chamber so as to penetrate the material; (d) evacuating gas from the sterilization chamber so as to evaporate the liquid at least partially; (e) introducing gaseous ozone into the sterilization chamber; (f) treating the material in a sterilization chamber for a sufficient time period so as to disinfect and sterilize the materials. The germicide potency of the method is markedly improved in comparison with the method where only compound O₃ is used alone or in combination with the germicide H₂O₂.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

A Method for Disinfecting and Sterilizing Microbial Contaminated Materials

The present invention relates to a method for disinfecting and sterilizing microbial contaminated materials, e.g. medical devices, instruments, waste from laboratories or hospitals comprising blood or blood components or other biological material under use of a combination of *in situ* generated peracetic acid and ozone.

For disinfecting of medical devices, working tools or infectious waste material in the medical field, several methods using ozone are suggested according to prior art. The main problem of known methods has been the long period for treating the material containing pathogenic agents such as bacteria and viruses. With a shorter treatment problems occurred with material containing biological liquids such as blood or components thereof (probably due to hemoglobin presence) because an amount of resistant spores survived. It was found that an improvement can be attained if a combination of ozone and peracetic acid was used in a humidified gaseous phase. Up to now peracetic acid has been used in liquid form for decontaminating medical wastes (US Patent 5,374,394) and the combination of peracetic acid and ozone as a disinfectant has never been used in liquid form. Another approach is described in the document JP-7-136236 wherein hot vapor is injected into a sterilization chamber. The use of a hot oxidizing agent can be irritating and dangerous in the case of a defect in the chamber.

The present invention is related to a method for disinfecting and sterilizing microbial contaminated or infectious materials, e.g. medical instruments, waste from medical laboratories or hospitals. The method comprises the steps:

- (a) loading the material for disinfecting and sterilizing into a pressure-proof sterilization chamber
- (b) introducing liquid hydrogen peroxide into the sterilization chamber so as to penetrate the material;
- (c) introducing liquid acetic acid into the sterilization chamber so as to penetrate the material;

- (d) evacuating gas from the sterilization chamber so as to evaporate the liquid at least partially;
 - (e) introducing gaseous ozone into the sterilization chamber;
 - (f) treating the material in the sterilization chamber for a sufficient time period so
- 5 as to disinfect and sterilize the materials

The method can be carried out in treatment devices as described in the patent documents EP-A- 0 664 715 and EP-A-0 761 237, which can be adapted to the method according to present invention.

The method of the present invention is carried out, as a rule, at a

10 temperature of about 15 °C to 35 °C and preferably at ambient temperature.

The hydrogen peroxide and the acetic acid can be introduced enclosed in two separate ampoules having such properties that after the evacuation of the chamber they are burst with release of the hydrogen peroxide and the acetic acid.

15 Alternatively the hydrogen peroxide and the acetic acid can be introduced enclosed in a container having at least two compartments being constructed in such a manner that the two compartments release their contents during the evacuation of the sterilizing chamber.

The method of the invention can be carried out in two or more cycles

20 by feeding at least two times hydrogen peroxide, the acetic acid and/or ozone into the sterilization chamber.

Alternatively the sterilization chamber can be equipped with storage and feeding means for feeding continuously hydrogen peroxide, acetic acid and ozone in several process cycles.

25 The molar ratio of acetic acid to ozone according to the invention is as a rule about 3/1 to 1/3 and the humidity in the chamber is at least 10 %.

The infectious waste material treated according to the method of the present invention can be solid material or a mixture of solid and liquid material. For disinfecting waste material safely, it is ground or broken up in other wise before, during or after the treatment with the disinfecting agents (hydrogen
5 peroxide and acetic acid).

Preferably a stable, storable and shippable disinfecting and sterilizing combination of agents is used together with ozone in the method of the invention. The combination possesses improved properties and comprises a two-part system: the first part consists of a mixture of acetic acid and water, and
10 the second consists of hydrogen peroxide and water. In this solution the peracetic acid is formed *in situ*. With this measure it is possible to take advantage of the outstanding disinfecting and sterilizing properties of peracetic acid without exposure to the danger of explosion and to the strong irritating properties to the skin and the eyes. The application of the peracetic acid
15 forming combination with ozone is carried out in a gaseous phase under reduced pressure (pressure < atmospheric pressure) and at ambient temperature. Hydrogen peroxide reacts with acetic acid to form of peracetic acid and water:



20 This is an equilibrium which can be shifted if the reaction takes place in the presence of a catalytic amount of acid (e.g. sulfuric acid). After termination of the disinfecting process the remaining ozone is destroyed, and, if necessary, the acid can be neutralized.

The present invention is illustrated with sterilizing tests carried out in
25 a sterilizing apparatus of the type "BOX O3" of Carbagas Aktiengesellschaft CH 3097 Liebefeld (Switzerland). This apparatus was designed for disinfecting infectious waste material. Normally in this device the waste material is ground , evacuated and treated one or several times with gaseous ozone in a sterilizing chamber. According to the present invention *in situ*-formed peracetic acid is
30 used additionally. For comparison, instead of peracetic acid only hydrogen

peroxide was used. For the test, pads loaded with infected sheep's blood were used. The results are listed in the table below.

Table: Number of viable and sporular bacterial forms destroyed in a sterilization chamber on a pad in the presence of blood of a sheep (100 µl per germ-carrier)

micro-organism	Number of cycles	Formation time	Compound 1	Compound 2	Reduction of micro-organisms
<i>S. aureus</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	3.87 log ₁₀
<i>S. aureus</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	7.90 log ₁₀
<i>S. aureus</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<8.11 log ₁₀
<i>E. hirae</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	2.23 log ₁₀
<i>E. hirae</i> +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥7 log ₁₀
<i>E. hirae</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<7.8 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<7.69 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	5	480 s	H ₂ O ₂	-	1.38 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<8 log ₁₀

<i>P. aeruginosa</i> +100µl of sheep's blood	5	480 s	H ₂ O ₂	-	1.47 log ₁₀
<i>P. aeruginosa</i> +100µl of sheep's blood	4	900 s	H ₂ O ₂	CH ₃ COOH	7 log ₁₀
<i>P. aeruginosa</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8.25 log ₁₀
<i>P. aeruginosa</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	8.47 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	2.47 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≤7 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≤7.6 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≤8 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	1.8 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8.04 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8.04 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<8 log ₁₀

<i>B.subtilis</i> spores +100µl of sheep's blood	3	480 s	H ₂ O ₂	-	3 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	5	600 s	H ₂ O ₂	CH ₃ COOH	≥3 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥6 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	4	900 s	H ₂ O ₂	CH ₃ COOH	≥6 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	4 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	5	480 s	H ₂ O ₂	-	<4 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥5 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥5 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥5 log ₁₀

The table shows that the germicide potency of the method according to the invention is markedly improved in comparison with the method where
5 only the germicide compound H₂O₂ is used in combination with O₃.

Claims

1. A method for disinfecting and sterilizing microbial contaminated or infectious materials, e.g. medical instruments, waste from medical laboratories or hospitals, the steps comprising:
 - 5 (a) loading the material for disinfecting and sterilizing into a vacuum-proof sterilization chamber
 - (b) introducing liquid hydrogen peroxide into the sterilization chamber so as to penetrate the material;
 - (c) introducing liquid acetic acid into the sterilization chamber so as to penetrate
10 the material;
 - (d) evacuating gas from the sterilization chamber so as to evaporate the liquid at least partially;
 - (e) introducing gaseous ozone into the sterilization chamber;
 - (f) treating the material in a sterilization chamber for a sufficient time period so
15 as to disinfect and sterilize the materials
2. The method of claim 1 wherein the steps (b) and (c) are carried out in any sequence under *in situ* generation of peracetic acid.
3. The method of claim 1 wherein the steps (b) and (c) are carried out simultaneously under *in situ* generation of peracetic acid.
- 20 4. The method according to one of the claims 1 to 3 wherein the method is carried out at a temperature of about 15 °C to 35 °C.
5. The method of claim 4 wherein the method is carried out at ambient temperature.
6. The method of claim 2 or 4 wherein the hydrogen peroxide and
25 the acetic acid of steps (b) and (c) are introduced in two separate ampoules having such properties that after the evacuation of step (d) they burst with release of their content.
7. The method of claim 3 or 4 wherein the hydrogen peroxide and the acetic acid of steps (b) and (c) are introduced in a container having at least
30 two compartments being constructed in such a manner that their contents are released is during the evacuation of step (d).

8. The method of claim 7 wherein the container has a compartment with a catalyst for shifting the equilibrium of the reaction equation



to the right, which catalyst is released at the same time as the hydrogen
5 peroxide and the acetic acid.

9. The method of claim 8 wherein the catalyst is sulfuric acid.

10. The method according to one of the claims 1 to 9 wherein in step (f) the ozone and the formed peracetic acid are present in the vapor phase.

11. The method according to one of the claims 1 to 10 wherein the
10 steps (b) to (e) are carried out at least two times.

12. The method according to one of the claims 1 to 11 wherein at the sterilization chamber feeding means for feeding hydrogen peroxide, acetic acid and ozone are disposed.

13. The method according to one of the claims 1 to 12 wherein the
15 molar ratio of acetic acid to ozone is 3/1 to 1/3.

14. The method according to one of the claims 1 to 13 wherein the humidity in step (e) or (f) is at least 10 %.

15. The method according to one of the claims 1 to 14 wherein the material is infectious waste in a solid or liquid state.

20 16. The method according to one of the claims 2 to 15 for treating contaminated or infectious waste wherein the waste is ground before, during or after step (b) or (c).

17. The method according to one of the claims 1 to 16 wherein step (e) is carried out at least twice.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/04723

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61L2/20 A61L11/00 //A61L101/10,A61L101/36

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 94 21120 A (ENVIRO MEDICAL SYSTEMS INC) 29 September 1994 (1994-09-29) page 8, line 21 - line 29 page 9, line 10 - line 17 page 9, line 27 -page 10, line 4 page 11, line 23 - line 27 ---	1-5, 7-12, 14-17
A	US 5 700 426 A (BARDAT ANNIE ET AL) 23 December 1997 (1997-12-23) column 2, line 35 - line 60 column 3, line 1 - line 3 claims 1,4,5 --- -/--	1-5, 7-12, 14-17



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

21 October 1999

Date of mailing of the international search report

03/11/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Menidjel, R

INTERNATIONAL SEARCH REPORT

Intern al Application No

PCT/EP 99/04723

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>US 5 567 444 A (HEI ROBERT D ET AL) 22 October 1996 (1996-10-22) column 1, line 54 -column 2, line 20 column 5, line 30 - line 59 column 10, line 10 - line 35 claims 1-7</p> <p style="text-align: center;">---</p>	1-17
A	<p>US 5 674 450 A (SWANZY JAMES ARCHIE ET AL) 7 October 1997 (1997-10-07) column 4, line 38 - line 52 column 5, line 9 - line 36</p> <p style="text-align: center;">-----</p>	1-17

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 99/04723

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9421120 A	29-09-1994	CA 2116281 A	18-09-1994
		CA 2158459 A	29-09-1994
		EP 0689378 A	03-01-1996
US 5700426 A	23-12-1997	FR 2673540 A	11-09-1992
		CA 2105778 A	09-09-1992
		DE 69223613 D	29-01-1998
		DE 69223613 T	28-05-1998
		EP 0574505 A	22-12-1993
		WO 9215337 A	17-09-1992
		JP 6505659 T	30-06-1994
US 5567444 A	22-10-1996	US 5484549 A	16-01-1996
		CA 2174277 A	23-03-1997
		DE 19630615 A	27-03-1997
		FR 2739041 A	28-03-1997
		GB 2305435 A, B	09-04-1997
		IT T0960753 A	13-03-1998
		JP 9125097 A	13-05-1997
		NL 1004029 C	29-12-1998
		NL 1004029 A	25-03-1997
		AU 681411 B	28-08-1997
		AU 6964094 A	22-03-1995
		CA 2169636 A	09-03-1995
		DE 69412838 D	01-10-1998
		DE 69412838 T	14-01-1999
		EP 0716686 A	19-06-1996
		JP 9501981 T	25-02-1997
		NZ 267362 A	24-02-1997
		WO 9506712 A	09-03-1995
US 5674450 A	07-10-1997	AU 704105 B	15-04-1999
		AU 1774995 A	09-11-1995
		AU 2242499 A	20-05-1999
		BR 9501831 A	05-03-1996
		CA 2147953 A	29-10-1995
		CN 1112446 A	29-11-1995
		EP 0679407 A	02-11-1995
		FI 952005 A	29-10-1995
		JP 8038583 A	13-02-1996
		US 5667753 A	16-09-1997
		US 5876666 A	02-03-1999
		US 5770739 A	23-06-1998
		US 5785934 A	28-07-1998
		SG 28258 A	01-04-1996
		ZA 9503409 A	28-10-1996

THIS PAGE BLANK (USPTO)